INFORMATION DISCLOSURE	Application Number		10566063
	Filing Date		2006-01-26
	First Named Inventor Alan N		Martin Birch
STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)	Art Unit		
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	1	3706810		1972-12-19	American Cyanamid Company	
	2	4599198		1986-07-08	Pfizer Inc.	
	3	4668769		1987-05-26	Dennis J. Hoover	
	4	4692522		1987-09-08	Merck & Co., Inc.	
	5	4720503		1988-01-19	Merck & Co., Inc.	
	6	4751231		1988-06-14	Merck & Co., Inc.	
	7	4786641		1988-11-22	Bayer Aktiengesellschaft	
	8	4794120		1988-12-27	Synthelabo	

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	9	5863903		1999-0	1-26	Novo Nordisk	A/S				
	10	5998463		1999-12	2-07	Pfizer Inc.					
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	1	200740	DD		А	1983-06-08	Karl Gewald et. al.				<b>✓</b>
	2	4445968	DE		A1	1996-06-27	Bayer AG				<b>V</b>
	3	0697403	EP		A1	1996-02-21	Sanofi				<b>V</b>
	4	0846464	EP		A2	1998-06-10	Pfizer Inc				

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5	0884050	EP	A1	1998-12-16	Novo Nordisk A/S	
6	0978279	EP	A1	2000-02-09	Pfizer Products Inc.	
7	1088824	EP	A2	2001-04-04	Pfizer Products Inc.	
8	1125580	EP	A2	2001-08-22	Pfizer Products Inc.	
9	1134213	EP	A2	2001-09-19	Pfizer Inc.	
10	1136071	EP	A2	2001-09-26	Pfizer Products Inc.	
11	1145717	EP	A2	2001-10-17	Pfizer Products Inc.	
12	1149580	EP	A1	2001-10-31	Pfizer Products Inc.	
13	1177791	EP	A2	2002-02-06	Pfizer Products Inc.	
14	1340500	EP	A1	2003-09-03	Pfizer Products Inc	
15	2081747	ES	A1	1996-03-01	Laboratories del Dr. Esteve, S.A.	<b>✓</b>

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16	04179949	JP	А	1992-06-26	Fuji Photo Film CO., Ltd.	<b>✓</b>
17	2001 089368	JP	А	2001-04-03	Tanabe Seiyaku Co., Ltd	<b>✓</b>
18	2001 206856	JP	А	2001-07-31	Pfizer Products Inc.	<b>V</b>
19	2001 247565	JP	А	2001-09-11	Pfizer Products Inc	<b>✓</b>
20	2004 196702	JP	А	2004-07-15	Yamanouchi Pharmaceuticals Co., Ltd.	<b>V</b>
21	93/25574	WO	A1	1993-12-23	Pfizer Inc.	
22	95/24391	WO	A1	1995-09-14	Novo Nordisk A/S	
23	96/39384	WO	A1	1996-12-12	Pfizer Inc.	
24	96/39385	WO	A1	1996-12-12	Pfizer Inc.	
25	97/09040	WO	A1	1997-03-13	Novo Nordisk	
26	97/31901	wo	A1	1997-09-04	Mikael Bols	

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27	97/45425	wo	A1	1997-12-04	Fujisawa Pharmaceutical Co., Ltd.	
28	98/27108	wo	A2	1998-06-25	Fujisawa Pharmaceutical Co., Ltd.	
29	98/40353	wo	A1	1998-09-17	Novo Nordisk A/S	
30	98/50359	wo	A1	1998-11-12	Novo Nordisk A/S	
31	99/26659	wo	A1	1999-06-03	Pfizer Products Inc.	
32	99/36393	wo	A1	1999-07-22	Tanabe Seiyaku Co., Ltd.	
33	00/42213	wo	A1	2000-07-20	The Research Foundation of State University of NY	
34	00/47206	wo	A1	2000-08-17	Novo Nordisk A/S	
35	01/05954	wo	A1	2001-01-25	Isis Pharmaceuticals Inc	
36	01/23347	wo	A1	2001-04-05	Novo Nordisk A/S	
37	01/32622	wo	A1	2001-05-10	AstraZeneca AB	

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38	01/32654	wo	A2	2001-05-10	Societe de Conseils de Recherches et D'App Sci	<b>✓</b>
39	01/52825	wo	A2	2001-07-26	Novartis AG	
40	01/68055	wo	A1	2001-09-20	Pfizer Products Inc.	
41	01/68092	wo	A2	2001-09-20	Pfizer Products Inc.	
42	01/68603	wo	A2	2001-09-20	Bristol-Myers Squibb Co.	
43	01/94300	wo	A1	2001-12-13	Aventis Pharma Deutshland GMBH	<b>✓</b>
44	01/96311	wo	A2	2001-12-20	Bristol-Myers Squibb Company	
45	01/96347	wo	A1	2001-12-20	Bristol-Myers Squibb Company	
46	02/20530	wo	A1	2002-03-14	AstraZeneca AB	
47	02/26714	wo	A1	2002-04-04	Takeda Chemical Industries	<b>✓</b>
48	02/34718	wo	A1	2002-05-02	Richter Gedeon Vegyeszeti Gyar RT	

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	49	02/36583	WO	A1	2002-05-10	Shionogi & Co., Ltd.		<b>✓</b>
	50	02/80844	WO	A2	2002-10-17	Genzyme Corporation		
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	1	CROCHET R.A., et al., J. Het. Chem., "Synthesis of Substituted Thieno[2,3-b] pyrroles," April 1974, 143-150, Vol. 11.						
	2	HARTMAN G.D., et al., "The Synthesis of 5-Alkylaminomethylthieno[2,3-b]Pyrrole-5-Sulfonamides," Heterocycles, 1989, 1943-1949, Vol. 29(10).						
	3	MCCORMACK J.G., et al., "Pharmacological Approaches to Inhibit Endogenous Glucose Production as a Means of Anti-diabetic Therapy," Curr. Pharmaceutical Design, 2001, 1451-1474, Vol. 7.						
	4	JAKOBSEN P., et al., "Iminosugars: Potential Inhibitors of Liver Glycogen Phosphorylase.," Bioorganic Med. Chem., 2001, 733-744, Vol. 9.						
	5	TREADWAY J.L., et al., "Glycogen phosphorlase inhibitors for treatment of type 2 diabetes mellitus," Exp. Opin. Invest. Drugs, 2001, 439-454, Vol. 10(3).						
	6	RATH V.L. et al., "Activation of Human Liver Glycogen Phosphorylase by Alteration of the Secondary Structure and Packing of the Catalytic Core," Mol. Cell, July 2000, 139-148, Vol. 6.						
	7	OIKONOMAKOS N.G., et al., "Allosteric inhibition of glycogen phosphorylase a by the potential antidiabetic drug 3-isopropyl 4-(2-chlorophenyl)-1,4-dihydro-1-ethyl-2-methyl-pyridine-3,5,6-tricarboxylate," Protein Sci., 1999, 1930-1945, Vol. 8.						

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Art Unit		
Examiner Name Not y		et assigned
Attorney Docket Number		101159-1P US

8	VENKATARANGAN P., et al., "Prediction of Ligand-Receptor Binding Thermodynamics by Free Energy Force Field Three-Dimensional Quantitative Structure-Activity Relationship Analysis: Applications to a Set of Glucose Analogue Inhibitors of Glycogen Phosphorylase," J. Med. Chem., 1999, 2169-2179, Vol. 42.	
9	HOOVER D.J., et al., "Indole-2-carboxamide Inhibitors of Human Liver Glycogen Phosphorylase," J. Med. Chem., 1998, 2934-2938, Vol. 41.	
10	MARTIN W.H., et al., "Discovery of a human liver glycogen phosphorylase inhibitor that lowers blood glucose in vivo," PNAS, Feb. 1998, 1776-1781, Vol. 95.	
11	SOMAN G., et al., "The Nature of the Binding Site for Aromatic Compounds in Glycogen Phosphorylase b," Biochem. J., 1975, 369-371, Vol. 147.	
12	SOMAN G., et al. "Aromatic Compounds as Allosteric Inhibitors of Glycogen Phosphorylase b," Biochimica et Biophysica Acta, 1974, 359-362, Vol. 358.	
13	ROSAUER K.G., et al., "Novel 3,4-Dihydroquinolin-2(1H)-one Inhibitors of Human Glycogen Phosphorylase a," Bioorganic & Medicinal Chemistry Letters, 2003, 4385-4388, Vol. 13.	
14	TEAGUE J. et al., "Mobilisation of Tissue Glycogen Following Inhibition of Glycogen Phosphorylase in fa/fa Rat," Diabetes, 53, Supp. 2, 2004, A365, 1521-P.	
15	VERTIGAN H. et al., "Impact of cell glycogen content on modulation of hepatocyte glucose metabolism by pharmacological agents," Diabetologia, 2004, A214, 589, Vol. 47, Supp.1.	
16	FONT M. et al. "Indoles and pyridazino[4,5-b]indoles as nonnucleoside analog inhibitors of HIV-1 reverse transcriptase", European Journal Med Chem, 1995, 963-71, Vol. 30.	
17	LIN T. et al. "Effects of Protein Binding and Experimental Disease States on Brain Uptake of Benzodiazepines in Rats", J Pharmacology & Eptl Therapeutics, 1990, 45-50, Vol. 253(1).	
18	VARNAVAS A. et al. "Quinolone Derivatives: Synthesis and Binding Evaluation on Cholecystokinin Receptors", Il Farmaco, 1996, 341-350, Vol. 51(5).	

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	19	PARSONS W H. et al. "Cholecystokinin Antagonists. Synthesis and Biological Evaluation of 3-Substitued Benzolactams", J Med Chem, 1989, 1681-5, Vol. 32.	
	20	FREEMAN S., et al., "Effect of Glucose on Rat and Human Liver Glycogen Phosphorylasea Activity and Potency of a Glycogen Phosphoylase Inhibitor," Diabetes, 52, Supp., 2003, A340, 1470-P.	
	21	TURNBULL A., et al., "Pharmacological Inhibition of Glycogen Phosphorylase (GP) Lowers Plasma Glucose in Rat Models of Type 2 Diabetes," Diabetes, 52, Supp., 2003, A343, 1485-P.	
	22	BIRCH A., et al., "Novel Thienopyrrole Glycogen Phosphorylase Inhibitors: In Vitro SAR and Crystallographic Studies," Poster, Cambridge Med Chem Symposium, Sept 2003.	
	23	HUDSON S., et al., "The effect of a glycogen phosphorylase inhibitor upon muscle fatigue in anaesthetised rats," J. Physiol., 2002, 52-53, Vol. 539.	
	24	VERTIGAN H. et al., "Impact of cell glycogen content on modulation of hepatocyte glucose metabolism by pharmacological agents", EASD Munich, 2004.	
	25	BARTLETT J. et al., "In Vitro and In Vivo Profile of Gpi688, a Novel, Potent Inhibitor of Glycogen Phosphorylase", ADA San Diego, 2005.	
	26	BENNETT S N L. et al., "Novel Orally Active Amino-indan Inhibitors of Glycogen Phosphorylase", Cambridge Med Chem Conference, Sept 2005. Poster EOM.	
	27	GREEN A R. et al., "The Glycogenic Action of Protein Targeting to Glycogen in Hepatocytes Involves Multiple Mechanisms Including Phosphorylase Inactivation and Glycogen Synthase Translocation", J Biol Chem, 2004, 46474-46482, Vol. 279(45).	
	28	ROBERTS P A. et al., "The temporal relationship between glycogen phosphorylase and activation of the pyruvate dehydrogenase complex during adrenaline infusion in resting canine skeletal muscle", J Physiology, 2002, 297-304, Vol. 545(1).	
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